Kinetic investigations into the possible cause of low serum histidine in rheumatoid arthritis

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SUMMARY To investigate the cause of low serum histidine in rheumatoid arthritis (RA) single oral and intravenous doses of L-histidine were administered to patients with active RA, and to an equal number of age and sex matched control subjects. In the first study 13 patients and their controls received a 100 mg kg⁻¹ dose of L-histidine as an aqueous slurry. Significant differences were seen in body weight, predose baseline serum histidine concentration, Cmax, t½, and area under curve, AUC₀-∞. In a second study six patients and six controls each received a 50 mg kg⁻¹ dose of L-histidine both orally and intravenously on two separate occasions. The patients with RA had a lower baseline serum histidine concentration, a lower volume of distribution, and a shorter plasma half life than the controls, but these differences were not statistically significant. No difference was seen in bioavailability or clearance. Low serum histidine in RA is unlikely to be due to malabsorption from the gut, uptake by abnormal gut flora, or increased metabolism.

The concentration of free serum histidine (i.e., histidine+trace metals–histidine complexes) is significantly decreased in patients with rheumatoid arthritis (RA). Gerber has shown that serum histidine levels are related to disease activity in patients with RA.

One possible cause of this hypohistidinaemia is defective absorption from the gut, perhaps due to excessive uptake or metabolism by atypical gut flora. There is some evidence to suggest that patients with chronic polyarthritis have an intestinal flora with the occurrence of atypical strains of Clostridium perfringens type A, for which tryptophan and histidine are essential amino acids. Bergstrom and Havermark showed that the jejunal mucosa of patients with RA had 5–10 fold greater histidine methyl esterase activity than a control group.5 Alternatively, there may be a fault in a specific amino acid transport system across the gut wall.

A second possible explanation for low serum histidine may be an increase in catabolism. Histidine can be degraded to glutamate by the urocanate pathway, but can also be directly decarboxylated to histamine. Histamine is a well recognised mediator of inflammation and therefore may be produced in large quantities in RA. In mammalian tissues the synthesis of histamine is catalysed by the enzyme histidine decarboxylase, which has a high specificity for histidine, as well as by non-specific aromatic L-amino acid decarboxylase found in several animal tissues, e.g., kidney, brain, and liver.

To shed some light on the cause of low serum histidine we have investigated the pharmacokinetics of L-histidine in two histidine loading studies in patients with RA, and in age and sex matched normal controls.

Patients and methods

All patients had classical or definite RA (American Rheumatism Association criteria) and active disease as judged by a physician and the presence of a raised C reactive protein level. Approval was obtained for the study from the local ethical committee for medical research. The patients continued to take their non-steroidal anti-inflammatory and analgesic therapy throughout the trial. They were all inpatients at the Regional Rheumatology Centre in Harrogate, with the exception of the three youngest
women in study 1. In both studies the patients were age and sex matched with normal healthy controls. The demographic data for these subjects are shown in Table 1.

Subjects starved from 10.00 pm the previous night but were allowed water and their usual medication. Dietary restrictions were lifted three hours after dosing. Smokers were asked to refrain from smoking until the end of the sampling on the study days. All drugs were administered with patients in the upright position.

STUDY 1
Thirteen normal healthy volunteers and 13 patients with active RA took a 100 mg kg\(^{-1}\) l-histidine dose as an aqueous slurry at 9.00 am on the study day. The l-histidine used was the free base (AnaLaR grade) obtained from British Drug Houses (BDH, Atherstone, UK). Venous blood samples (10 ml) were taken by means of an indwelling cannula at times 0, 15, 30, 40, 50, 60, 70, 80, 90 minutes and 2, 3, 4, 6, and 8 hours relative to the dose. The samples were allowed to clot in plain glass tubes at room temperature and then centrifuged at 550 g for 10 minutes within two hours of clotting. The serum was pipetted into a separate plastic tube and stored at \(-20^\circ\)C before analysis.

STUDY 2
Six female patients with RA and six healthy female controls each received a 50 mg kg\(^{-1}\) body weight oral histidine bolus (as an aqueous slurry), and 50 mg kg\(^{-1}\) body weight intravenous (IV) solution. The two doses were separated by seven to eight days. In each group the dosing order was random.

Each subject was given a single oral dose of mepyramine (100 mg) before each dose of histidine to prevent any local inflammatory reaction at the site of entry of the infusion. It was given with the oral dose also to standardise trial conditions on both occasions. The intravenous solution (35 mg ml\(^{-1}\) l-histidine) was prepared and packed by the drug manufacturing department at St James's Hospital, Leeds. The solution was administered at a mean rate of 154 ml h\(^{-1}\) (range 99–223 ml h\(^{-1}\)) using a paediatric giving set which allowed an accurate amount to be given, though this resulted in a long infusion.

An intravenous cannula from which blood samples were taken was inserted into a forearm vein contralateral to the infusion. After the oral dose, blood samples (10 ml) were taken at 0, 10, 20, 30, 40, 50, 60, 75, 90, 105, 120, 150, 180, 240, 300, 360, and 480 minutes. For the IV dose, samples were taken at 10 minute intervals until the end of the infusion, and then at 5, 10, 15, 25, 35, 45, 60, 75, 90, 105, 120, 150, 180, 210, 240, 270, and 300 minutes. Samples were treated and stored as described for study 1.

LABORATORY ASSESSMENTS
Total serum histidine concentration was assayed by a modification of the method of Gerber\(^8\) employing a Perkin Elmer LS3 spectrofluorimeter. A six point calibration (10–200 mg l\(^{-1}\)) was employed with precision ranging from 8.3% to 0.4% (mean 2.6%) and accuracy from 93.0% to 101.8%. The average recovery was 95 (SEM 2.7%). Quality control samples were included in every analytical run, with the run being rejected if the results for the control samples were 10% more or less than the previously determined value. Further details relating to the assay and sample storage are given elsewhere.\(^9\)

PHARMACOKINETIC ANALYSIS
Model dependent
Individual data sets were best fitted, using

<table>
<thead>
<tr>
<th>Parameter</th>
<th>RA</th>
<th>Control</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex (F:M)</td>
<td>9:4</td>
<td>9:4</td>
<td>—</td>
</tr>
<tr>
<td>Age (years)</td>
<td>40 (4)</td>
<td>39 (4)</td>
<td>NS</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>55 (3)</td>
<td>64 (2)</td>
<td>&lt;0.02</td>
</tr>
<tr>
<td>Dose (g)</td>
<td>5.5 (0.27)</td>
<td>6.4 (0.19)</td>
<td>&lt;0.02</td>
</tr>
<tr>
<td>Initial, baseline histidine (mg l(^{-1}))</td>
<td>15.0 (0.7)</td>
<td>18.0 (0.7)</td>
<td>&lt;0.007</td>
</tr>
<tr>
<td>C(_{max}) (mg l(^{-1}))</td>
<td>134 (8)</td>
<td>158 (5)</td>
<td>&lt;0.016</td>
</tr>
<tr>
<td>T(_{max}) (h)</td>
<td>0.72 (0.05)</td>
<td>0.72 (0.04)</td>
<td>NS</td>
</tr>
<tr>
<td>t(_{1/2}) (h)</td>
<td>1.18 (0.16)</td>
<td>1.69 (0.19)</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>AUC(_{0-\infty}) (mg l(^{-1}) h)</td>
<td>312 (44)</td>
<td>477 (49)</td>
<td>&lt;0.02</td>
</tr>
</tbody>
</table>

NS=not significant. p>0.05.

C\(_{max}\)=maximum serum histidine concentration; T\(_{max}\)=time to achieve C\(_{max}\); t\(_{1/2}\)=serum histidine half life; AUC=area under the serum concentration versus time curve.
ELSOMS,\textsuperscript{10} to a one compartment model having first order absorption and first order elimination superimposed on a constant rate input equal to the rate of output under basal conditions. If basal serum level=$s_0$ (mg l\textsuperscript{-1}), volume=$V$ (l), and elimination rate constant=$ke$ (h\textsuperscript{-1}) then the rate of output=$s_0\times V\times ke$=rate of input=(mg h\textsuperscript{-1}). When dosed with the drug, input=$s_0\times V\times ke+A$, where for the IV dose $A$=dose/infusion time and for the oral dose $A=f\times\text{dose}\times ka$, where $f$=fraction of dose remaining to be absorbed and $ka$=absorption rate constant. For the oral dose $A=0$ during the time lag before absorption starts. Output=$s\times V\times ke$, where $s$=serum histidine level.

All data sets were fitted to this model and the best values of $s_0$, $V$, and $ke$ were found together with $ka$ for oral doses. All sets gave an acceptable fit, though $s_0$ differed in some cases from the measured value. Clearance was calculated as $ke\times V$ (1 h\textsuperscript{-1}). Basal flux (or turnover) ($F_I$) was $ke\times V\times s_0$ (mg h\textsuperscript{-1}) from IV data. Bioavailability (%) was calculated as $100\times Cl$ (IV)/Cl (oral) and $t_{1/2}$ as $1\ln 2/ke$, where Cl=total body clearance.

Model independent

Computer modelling showed that the data most closely fitted a one compartment model with first order elimination. Maximum serum histidine concentration ($C_{\text{max}}$) and time to achieve $C_{\text{max}}$ ($T_{\text{max}}$) were observed directly from the data. (In the case of IV dosing $C_{\text{max}}$ was the concentration at the completion of the infusion.) The serum histidine half life ($t_{1/2}$) was obtained by log linear least squares regression analysis of data points following the peak concentration, after subtraction of the baseline histidine concentration from each result.

The area under the serum concentration versus time plot (AUC\textsubscript{0-\infty}) was calculated using the trapezoidal rule. As the AUC was required to determine bioavailability the initial baseline histidine concentration was subtracted from the total concentration at each time point. The apparent volume of distribution ($V_D$) was calculated from

$$V_D = D \times t_{1/2} \over \text{AUC}_{0-\infty} \times \ln 2$$

where $D$=dose and complete drug bioavailability after an IV dose is assumed. Total body clearance (Cl) was determined from $D$/AUC\textsubscript{0-\infty}.

**Statistical analysis**

An unpaired two sample $t$ test (two tailed) was used to compare the pharmacokinetic data for patients and controls in both studies.

**Results**

**STUDY 1**

The kinetic results are summarised in Table 1. Predose serum histidine concentrations were significantly lower in the patients. When patients and controls were compared the patients showed a significantly lower $C_{\text{max}}$, shorter $t_{1/2}$, and smaller AUC\textsubscript{0-\infty}. Dose and hence body weight correlated significantly ($p<0.001$) with $C_{\text{max}}$ and AUC\textsubscript{0-\infty} but not with $t_{1/2}$ in both patients and controls.

**STUDY 2**

In this study patients were matched to controls not only for age and sex but also for body weight, and hence for dose. Although the absolute results generated by the model dependent (Table 2) and model independent (Table 3) analyses were not identical, the trends in the differences between patients and controls were the same. There were no

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Oral dose</th>
<th>IV dose</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Controls</td>
<td>RA</td>
</tr>
<tr>
<td></td>
<td>$3.07$ (0-40)</td>
<td>$3.89$ (1-61)</td>
</tr>
<tr>
<td></td>
<td>$1.04$ (0-27)</td>
<td>$1.19$ (0-20)</td>
</tr>
<tr>
<td>$V$ (l)</td>
<td>$21.3$ (3-2)</td>
<td>$25.8$ (3-0)</td>
</tr>
<tr>
<td>$s_0$ (mg l\textsuperscript{-1})</td>
<td>$20.2$ (0-9)</td>
<td>$17.3$ (1-0)</td>
</tr>
</tbody>
</table>
| Calculated parameters | \begin{tabular}{l} 
$29.2$ (5-1) \hfill $29.1$ (4-1) \hfill $23.5$ (2-7) \hfill $25.9$ (3-0) \\
$- $ \hfill $- $ \hfill $485$ (90) \hfill $494$ (61) \\
$0.88$ (0-18) \hfill $0.68$ (0-13) \hfill $0.89$ (0-15) \hfill $0.59$ (0-10) \\
$86.3$ (8-3) \hfill $97.8$ (18-7) \hfill $- $ \hfill $- $
\end{tabular} |

*Values are mean (SEM).
The table below summarizes the model independent pharmacokinetic parameters in study 2:

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Oral dose</th>
<th>IV dose</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Controls (mg l⁻¹)</td>
<td>RA (mg l⁻¹)</td>
</tr>
<tr>
<td>Baseline histidine</td>
<td>15.4 (0.9)</td>
<td>14.8 (1.7)</td>
</tr>
<tr>
<td>Cmax (mg l⁻¹)</td>
<td>73.1 (6.6)</td>
<td>90.1 (6.9)</td>
</tr>
<tr>
<td>t½ (h)</td>
<td>1.14 (0.13)</td>
<td>1.02 (0.16)</td>
</tr>
<tr>
<td>AUC0-∞ (mg l⁻¹ h)</td>
<td>142 (18)</td>
<td>178 (39)</td>
</tr>
<tr>
<td>VD (l kg⁻¹)</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Cl (l h⁻¹)</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Bioavailability (%)</td>
<td>80 (9.9)</td>
<td>91 (5.1)</td>
</tr>
</tbody>
</table>

*Values are mean (SEM).

Statistically significant differences between patients and controls for any kinetic parameter, using either the model dependent or model independent results. The improved matching showed similar AUC0-∞ results in patients and controls after both oral and IV doses, but the bioavailability was found to be quite variable (52–111%) (Tables 2 and 3). The serum t½ was shorter and the Cmax higher in patients than in the controls after oral and IV doses. VD was notably lower in five of the six patients, while Cl showed no difference. There was no significant correlation between dose and any kinetic parameter.

**Discussion**

Some of the kinetic differences between patients with RA and normal controls observed in the first study may be attributed to the differences in body weight (e.g. AUC0-∞). Although the dose was titrated against body weight, this could have confounded the results from this study as the differences might be explained on the basis of the different doses. In the second study the two groups were matched for weight in addition to age and sex, and hence the trends observed in the results might be more reliably attributed to disease. Differences in baseline histidine and t½ seen in the first study were maintained in the second study, but with the small number of patients in the latter it was not surprising that these failed to achieve statistical significance.

Bioavailability was variable though slightly greater in RA, suggesting an absence of abnormal histidine absorption in RA. These results do not exclude the possibility that patients with RA may have difficulty digesting protein such that low net histidine absorption arises. Concentrations of histidine in the bowel will clearly usually be lower than those achieved in these studies.

The major kinetic difference observed was a smaller VD, resulting in a shorter serum t½ in patients. The smaller volume cleared at the same rate would take less time, and this is compatible with identical clearance results in the two groups. The differences in VD are not readily explained, but differences in non-specific histidine binding may be considered. In RA there are changes in protein concentrations, particularly the acute phase proteins. Some basic drugs, e.g., propranolol, are known to bind to α1 acid glycoprotein, and so a basic amino acid such as histidine may also bind to an acidic acute phase protein. Changes in protein conformation may also occur in RA, which could affect the binding capacity. This point may be further complicated by the presence of bound anti-inflammatory drugs in the patients. Alternatively, there may be a decrease in tissue binding of histidine arising from displacement by drugs, though the VD results are small in both groups, suggesting limited tissue uptake. Increased urine concentrations of urocanic acid have been observed in RA, and these were considered to be related to pyridoxine deficiency (pyridoxal phosphate is an essential cofactor for urocanase) rather than to folate status.11 Our results indicate that this failure of patients to degrade histidine fully does not affect its clearance.

There remains the possibility that if a different amino acid was used in a similar loading study that similar results might be obtained in the absence of a difference in baseline levels. Future studies should therefore include a loading study similar to study 2 using an amino acid which shows normal serum levels in RA. To be certain in the selection of the latter, however, the levels of other amino acids in RA require confirmation as previous investigations have only agreed on abnormally low serum histidine.12–15

It can be concluded that there is a trend towards a reduction in the distribution volume of a histidine load in patients with RA. This occurs in conjunction with a reduction in serum half life but no change in clearance. There is no evidence for abnormal
histidine absorption or increased metabolism in RA, but the precise cause of low serum histidine remains obscure.

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References

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